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SEARCH REQUEST FORM

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Scientific and Technical Information Center

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Requester's Full Name:	Sabiha C	250	Examiner #:	74141	Date: 2	15/0.	2
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If more than one search is			e searches in o	rder of nee	ed. ******	M/S	*****
Please provide a detailed statement							
Include the elected species or struc utility of the invention. Define any known. Please attach a copy of the	y terms that may have cover sheet, pertinen	a special me t claims, and	eaning. Give exampl abstract.	les or relevant	citations, a	uthors, etc,	if · ·
Title of Invention: & /	trogenie	Co-	for as	Anti	ange	ögen	ic_
Inventors (please provide full na	$_{\text{mes}}$): $D'A$.	mato	et al.	•	U		age of the second
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=> fil reg FILE 'REGISTRY' ENTERED AT 15:28:37 ON 19 FEB 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 American Chemical Society (ACS)

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STRUCTURE FILE UPDATES: DICTIONARY FILE UPDATES: 18 FEB 2002 HIGHEST RN 393508-26-4

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

The P indicator for Preparations was not generated for all of the CAS Registry Numbers that were added to the H/Z/CA/CAplus files between 12/27/01 and 1/23/02. Use of the P indicator in online and SDI searches during this period, either directly appended to a CAS Registry Number or by qualifying an L-number with /P, may have yielded incomplete results. As of 1/23/02, the situation has been resolved. Also, note that searches conducted using the PREP role indicator were not affected.

Customers running searches and/or SDIs in the $\rm H/Z/CA/CAplus$ files incorporating CAS Registry Numbers with the P indicator between 12/27/01 and 1/23/02, are encouraged to re-run these strategies. Contact the CAS Help Desk at 1-800-848-6533 in North America or 1-614-447-3698, worldwide, or send an e-mail to help@cas.org for further assistance or to receive a credit for any duplicate searches.

=> d sta que 129

L13

32 SEA FILE=REGISTRY ABB=ON PLU=ON (1035-77-4/BI OR 117048-59-6/ BI OR 1236-72-2/BI OR 15833-07-5/BI OR 16205-32-6/BI OR 165619-07-8/BI OR 165619-10-3/BI OR 165619-11-4/BI OR 165619-22 -7/BI OR 165619-23-8/BI OR 192062-02-5/BI OR 192062-12-7/BI OR 192062-13-8/BI OR 192062-14-9/BI OR 192062-15-0/BI OR 192062-20 -7/BI OR 22415-44-7/BI OR 26788-23-8/BI OR 26890-04-0/BI OR 302799-37-7/BI OR 362-07-2/BI OR 362-08-3/BI OR 383414-35-5/BI OR 50-27-1/BI OR 50-28-2/BI OR 518-28-5/BI OR 53-16-7/BI OR 56-53-1/BI OR 57-63-6/BI OR 5976-67-0/BI OR 64-86-8/BI OR 95041-90-0/BI)

27 SEA FILE=REGISTRY ABB=ON PLU=ON L13 AND C5-C6-C6-C6/ES L14L16

Ak-Cy C=C-G3 @38 39 40 Ak-C=C $N \sim N \sim N$ $C \equiv C - G3$ @30 31 32 @33 34 35 @36 @37 @41 42 43 C≡C 051 52 Ak-G4 055 56 Ak—Cy—C=C @50 47 48 49 cy-c=c0— G3 053 54 044 45 46

VAR G1=H/X/CN/AK/OH/22/NH2/24/26 VAR G2=30/CN/33/38/41/44/50/51/53/55/62/65/66 VAR G3=AK/CY/36 VAR G4=OH/NH2/X/58VAR G5=AK/CY/36-66 37-68 NODE ATTRIBUTES: CONNECT IS M1 RC AT CONNECT IS M1 RC AT CONNECT IS M1 RC AT 15 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

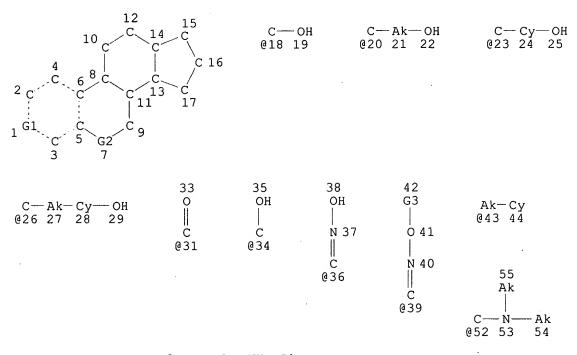
GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 67

STEREO ATTRIBUTES: NONE

L18 394 SEA FILE=REGISTRY CSS FUL L16

L19 · STR



VAR G1=C/18/20/23/26
VAR G2=C/31/34/36/39/45/47/49/52
VAR G3=AK/CY/43
NODE ATTRIBUTES:
CONNECT IS M1 RC AT 2
CONNECT IS M1 RC AT 3
CONNECT IS M1 RC AT 4
CONNECT IS M1 RC AT 14
CONNECT IS M1 RC AT 15
DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS AIOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RSPEC 1

NUMBER OF NODES IS 53

STEREO ATTRIBUTES: NONE

L21 152 SEA FILE=REGISTRY SUB=L18 CSS FUL L19

L22 STR

VAR G1=C/25/19/22/27/33/37/39/42/46 VAR G2=AK/CY/31 NODE ATTRIBUTES: CONNECT IS M1 RC AT CONNECT IS M1 RC AT CONNECT IS M1 RC AT 3 CONNECT IS M1 RC AT CONNECT IS M1 RC AT CONNECT IS M1 RC AT DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RSPEC 1

NUMBER OF NODES IS 46

STEREO ATTRIBUTES: NONE

L24 88 SEA FILE=REGISTRY SUB=L21 CSS FUL L22
L25 13 SEA FILE=REGISTRY ABB=ON PLU=ON L14

L25 13 SEA FILE=REGISTRY ABB=ON PLU=ON L14 AND L24 L27 75 SEA FILE=REGISTRY ABB=ON PLU=ON L24 NOT L25

L28 68 SEA FILE=REGISTRY ABB=ON PLU=ON L27 NOT (IDS/CI OR (D OR

T)/ELS OR 11C# OR 13C# OR 14C#)

L29 81 SEA FILE=REGISTRY ABB=ON PLU=ON (L25 OR L28)

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(FILE 'HOME' ENTERED AT 14:20:04 ON 19 FEB 2002) SET COST OFF

FILE 'HCAPLUS' ENTERED AT 14:20:15 ON 19 FEB 2002

E DAMATO/AU E D AMATO/AU

L1 65 S E74, E77, E84, E86, E87

E AMATO/AU

L2 2 S E36

L3 28 S E133, E135, E140, E141

E VARMA R/AU

L4 124 S E3, E10, E11, E12

L5 133 S E45-E47

E HAUGWITZ R/AU

L6 147 S E3-E11

E CUSHMAN M/AU

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L7
           201 S E3-E6, E8, E9, E10
            685 S L1-L7
L8
L9
             14 S L8 AND ?ESTROGEN?
L10
             42 S L8 AND ?ANGIO?
L11
              4 S L9 AND L10
              1 S L11 AND ESTROGEN?/TI
L12
                SEL RN
     FILE 'REGISTRY' ENTERED AT 14:24:13 ON 19 FEB 2002
             32 S E1-E32
L13
             27 S L13 AND C5-C6-C6/ES
L14
                                                          Species
L15
              1 S L14 AND C21H28O2
                E C21H28O2/MF
L16
                STR
L17
              9 S L16 CSS SAM
L18
            394 S L16 CSS FUL
                SAV L18 QAZ899/A
L19
                STR
L20
             5 S L19 CSS SAM SUB=L18
            152 S L19 CSS FUL SUB=L18
L21
                SAV L21 QAZ899A/A
L22
                STR
L23
             4 S L22 CSS SAM SUB=L21
L24
             88 S L22 CSS FUL SUB=L21
                SAV L24 QAZ899B/A
L25
             13 S L14 AND L24
             14 S L14 NOT L25
L26
             75 S L24 NOT L25
L27
L28
             68 S L27 NOT (IDS/CI OR (D OR T)/ELS OR 11C# OR 13C# OR 14C#)
             81 S L25, L28
L29
             80 S L29 NOT L15
L30
                SAV L30 QAZ899C/A
     FILE 'HCAOLD' ENTERED AT 15:15:35 ON 19 FEB 2002
L31
             0 S L15
L32
             72 S L30
     FILE 'HCAPLUS' ENTERED AT 15:15:58 ON 19 FEB 2002
                                                               - Speues
              2 S L15
L33
L34
              2 S L8 AND L33
L35
            489 S L30
                E ANGIOGENESIS/CT
                E E3+ALL
L36
           5343 S E5+NT
                E E10+ALL
          1959 S E4+NT
L37
L38
          73654 S E3+NT
                E E3+ALL
         77026 S E4
L39
L40
         165292 S E501+NT
L41
             13 S L35 AND L36
             83 S L35 AND L37-L41
L42
             38 S L35 AND ANGIOGENES?
L43
L44
              3 S L35 AND ANTIANGIOGENES?
L45
             15 S L35 AND ANTIANGIO?
             39 S L41, L43-L45
L46
L47
             33 S L42 AND L46
             49 S L35 AND P/DT
L48
            358 S L35 AND ?ESTROGEN?
L49
             67 S L49 AND L41-L48
L50
             19 S L50 AND L46
L51
L52
             30 S L35 AND ANGIO?/CW
L53
             29 S L52 AND L41-L48, L50, L51
L54
             30 S L34, L53
L55
             30 S L54 AND L33-L54
L56
              2 S L55 AND L34
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L57
             28 S L55 NOT L56
L58
             81 S L41-L48, L50-L55 NOT L57
L59
             64 S L30 (L) THU/RL
             36 S L59 AND L58
L60
             16 S L60 AND (1 OR 63)/SC
L61
             13 S L60 AND (1 OR 63)/SX
L62
             24 S L61, L62
L63
             24 S L59 AND ?ANGIO?
L64
             24 S L64 AND L33-L63
L65
             23 S L65 NOT 27/SC
L66
             24 S L33, L34, L66 AND L33-L66
L67
                SEL HIT RN
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FILE 'REGISTRY' ENTERED AT 15:28:10 ON 19 FEB 2002

L68 17 S E1-E17 L69 16 S L68 NOT L15

FILE 'REGISTRY' ENTERED AT 15:28:37 ON 19 FEB 2002

- hit compounds for L67, 1-24

=> d ide can 115

ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS L15

RN 165619-11-4 REGISTRY

Estra-1,3,5(10)-triene-3,17-diol, 2-(1E)-1-propenyl-, (17.beta.)- (9CI) CN (CA INDEX NAME)

OTHER CA INDEX NAMES:

Estra-1,3,5(10)-triene-3,17-diol, 2-(1-propenyl)-, [2(E),17.beta.]-

OTHER NAMES:

CN NSC 667047

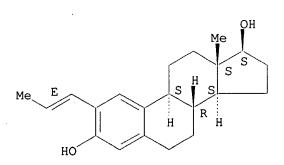
STEREOSEARCH FS

MF C21 H28 O2

SR

CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL LC STN Files:

Absolute stereochemistry. Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 136:64669 REFERENCE

123:112496 REFERENCE 2:

=> fil hcaplus FILE 'HCAPLUS' ENTERED AT 15:28:54 ON 19 FEB 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 18 Feb 2002 VOL 136 ISS 8 FILE LAST UPDATED: 17 Feb 2002 (20020217/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d 170 bib abs hitrn tot

L70 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2002 ACS

AN 2002:11127 HCAPLUS

DN 136:64669

TI Estrogenic compounds as antiangiogenic agents

IN D'Amato, Robert J.; Varma, Ravi K.; Haugwitz,
Rudiger G.; Cushman, Mark

Ι

PA USA

SO U.S. Pat. Appl. Publ., 14 pp., Cont. of U.S. Ser. No. 154,322, abandoned. CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

17111	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI PRAI	US 2002002294 US 1997-59916 US 1998-154322	A1 P B1	20020103 19970924 19980916	US 2001-899702	20010705
OS GI	MARPAT 136:64669				

$$R^2$$
 R^3
 Me
 R^4
 R^4

2-Methoxyestradiol derivs., such as I [R1, R3 = H, C1, Br, I, F, CN, alkyl, OH, CH2OH, NH2, alkylamino; R2 = N3, CN, C.tplbond.CR, C=CHR, RCH=CH2, C.tplbond.CH, OR, R-R1, OR-R1 (R = alkyl, R1 = OH, NH2, C1, Br, I, F, CF3); Z = CH, COH, CR2-OH (R2 = alkyl, aralkyl); Z' = CH2, CO, CH(OH); C=NOH, C=NOR5, CHC.tplbond.N, CHNR5R5 (R5 = H, alkyl, aralkyl)], were used for treating mammalian disease characterized by undesirable angiogenesis. Thus, 2-methoxyestradiol (II) showed inhibition of tubulin polymn. (IC50 = 3.6.+-.0.4 .mu.M), inhibition of colchicine binding to tubulin (1.9.+-.0.2 .mu.M) and antitumor activity against breast, CNS, melanoma, ovarian tumor cell assay in vitro.

IT 362-07-2, NSC 659853 362-08-3, 2-Methoxyestrone
22415-44-7 165619-07-8, NSC 671043 165619-11-4
, NSC 667047 192062-02-5, NSC 682429 192062-12-7, NSC
679431 192062-13-8, NSC 681684 192062-14-9, NSC 680185
192062-15-0, NSC 681683 192062-20-7, NSC 683125
302799-37-7, NSC 683688 383414-35-5, NSC 678473
RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(estrogenic compds. as antiangiogenic agents)

L70 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2002 ACS

AN 1995:592276 HCAPLUS

DN 123:112496

- TI Synthesis, Antitubulin and Antimitotic Activity, and Cytotoxicity of Analogs of 2-Methoxyestradiol, an Endogenous Mammalian Metabolite of Estradiol That Inhibits Tubulin Polymerization by Binding to the Colchicine Binding Site
- AU Cushman, Mark; He, Hu-Ming; Katzenellenbogen, John A.; Lin, Chii M.; Hamel, Ernest
- CS Department of Medicinal Chemistry, Purdue University, West Lafayette, IN, 47907, USA
- SO J. Med. Chem. (1995), 38(12), 2041-9 CODEN: JMCMAR; ISSN: 0022-2623
- DT Journal
- LA English
- In order to define the structural parameters assocd. with the antitubulin AΒ activity and cytotoxicity of 2-methoxyestradiol, a mammalian metabolite of estradiol, an array of analogs was synthesized and evaluated. The potencies of the new congeners as inhibitors of tubulin polymn. and colchicine binding were detd. using tubulin purified from bovine brain, and the cytotoxicities of the new compds. were studied in a variety of cancer cell cultures. Maximum antitubulin activity was obsd. in estradiols having unbranched chain substituents at the 2-position with three non-hydrogen atoms. 2-Ethoxyestradiol and 2-((E)-1propenyl)estradiol were substantially more potent than 2-methoxyestradiol The tubulin polymn. inhibitors in this series displayed significantly higher cytotoxicities in the MDA-MB-435 breast cancer cell line than in the other cell lines studied. The potencies of the analogs as cytotoxic and antimitotic agents in cancer cell cultures correlated with their potencies as inhibitors of tubulin polymn., supporting the hypothesis that inhibition of tubulin polymn. is the mechanism of the cytotoxic action of 2-methoxyestradiol and its congeners. Several of the more potent analogs were tested in an estrogen receptor binding assay, and their affinities relative to estradiol were found to be very
- IT 165619-11-4P 165619-12-5P 165619-13-6P

RL: BAC (Biological activity or effector, except adverse); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (synthesis, antitubulin and antimitotic activity, and cytotoxicity of analogs of methoxyestradiol)

IT 362-07-2P 165619-07-8P 165619-08-9P 165619-09-0P

RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (synthesis, antitubulin and antimitotic activity, and cytotoxicity of analogs of methoxyestradiol)

=> d 171 bib abs hitrn tot

- L71 ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:798040 HCAPLUS
- DN 135:339222
- TI Inhibition of abnormal cell proliferation with camptothecin or a derivative, analog, metabolite, or prodrug thereof, and combinations

```
including camptothecin
ΙN
     Rubinfeld, Joseph
PA
     Supergen, Inc., USA
ŞO
     PCT Int. Appl., 38 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                      KIND DATE
                                           APPLICATION NO.
     PATENT NO.
                                                            DATE
                                           -----
     WO 2001080843
                       Α2
                            20011101
                                           WO 2001-US12848 20010419
PΙ
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
             HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
             LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
             RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
             VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRAI US 2000-553710
                      A1
                            20000420
     A method for treating diseases assocd. with abnormal cell proliferation
     comprises delivering to a patient in need of treatment a compd. selected
     from 20(S)-comptothecin, an analog of 20(S)-comptothecin, a deriv. of
     20(S)-camptothecin, a prodrug of 20(S)-camptothecin, and pharmaceutically
     active metabolite of 20(S)-camptothecin, in combination with an effective
     amt. of one or more agents selected form the group consisting of
     alkylating agent, antibiotic agent, antimetabolic agent, hormonal agent,
     plant-derived agent, anti-angiogenesis agent and biol. agent.
     The method can be used to treat benign tumors, malignant or metastatic
     tumors, leukemia and diseases assocd. with abnormal angiogenesis
ΙT
     362-07-2, 2-Methoxyestradiol
     RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (camptothecin or deriv., analog, metabolite, or prodrug thereof for
        inhibition of abnormal cell proliferation, and combinations including
        camptothecin)
L71
     ANSWER 2 OF 22 HCAPLUS COPYRIGHT 2002 ACS
     2001:769051 HCAPLUS
ΑN
DN
     136:112738
     2-Methoxyestradiol: A novel endogenous chemotherapeutic and anti-
ΤI
     angiogenic agent
     Pribluda, Victor S.; LaVallee, Theresa M.; Green, Shawn J.
ΑU
CS
     EntreMed, Rockville, MD, USA
     New Angiotherapy (2002), 387-407. Editor(s): Fan, Tai-Ping D.; Kohn,
SO
     Elise C. Publisher: Humana Press Inc., Totowa, N. J.
     CODEN: 69BYCE
DT
     Conference; General Review
LA
     English
     A review which discusses the properties of 2-methoxyestradiol, which is a
AB
     part of a new generation of anti-cancer agents. The 2-methoxyestradiol is
     now recognized as a potent endogenous antimitogen and has the ability to
     selectively inhibit the growth of rapidly proliferating endothelial cells.
     Its candidacy as a viable antimitotic drug is supported by exptl. evidence
     showing that 2-methoxyestradiol is orally-active, and at therapeutic doses
     shows minimal, if any, toxicity.
     362-07-2, 2-Methoxyestradiol
IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use);
     BIOL (Biological study); USES (Uses)
        (methoxyestradiol as chemotherapeutic and anti-angiogenic
        agent)
              THERE ARE 108 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
        108
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ALL CITATIONS AVAILABLE IN THE RE FORMAT

```
L71
    ANSWER 3 OF 22 HCAPLUS COPYRIGHT 2002 ACS
ΑN
     2001:746919 HCAPLUS
DN
     136:31843
ΤI
     Angiogenetic and anti-angiogenetic effects of
     estradiol and its metabolites
     Mueck, A. O.; Seeger, H.; Lippert, C.; Wallwiener, D.
ΑU
CS
     Section of Endocrinology and Menopause, University Hospital, Tuebingen, 72
     076, Germany
     Journal of Clinical and Basic Cardiology (2001), 4(2), 153-155
SO
     CODEN: JCBCFT; ISSN: 1561-2775
     Krause & Pachernegg GmbH
PB
DΤ
     Journal
LA
     English
     Atherosclerotic plaques in later stages exhibit marked presence of new
AΒ
     micro vessels. Thus angiogenesis may be important for the
     development of atherosclerotic plaques and long-term anti-
     angiogenetic therapy may present an effective new
     anti-atherosclerotic approach. 2-Methoxyestradiol, an endogenous
     estradiol metabolite, has already been shown to be an effective anti-
     angiogenetic substance. In the present study 14 endogenous
     estradiol metabolites were tested on their angiogenetic and
     anti-angiogenetic properties and compared to the effect of their
     parent substance, 17.beta.-estradiol. Endothelial cells from human
     umbilical veins were used for the expts. 17.beta.-Estradiol showed a
     biphasic reaction on the proliferation of vascular endothelial cells.
     low concn. it stimulated and at high concns. it inhibited cell growth.
     The same pattern was obsd. for the hydroxylated A-ring metabolites.
     Methylation of these metabolites, however, completely abrogated the
     anti-proliferative effect at high concns., except for the metabolite
     2-hydroxyestradiol. For the D-ring metabolites no marked changes were
     obsd. These results indicate that in addn. to 2-methoxyestradiol other
     endogenous estradiol metabolites are potent anti-angiogenetic
     substances at high dosages. Since some of these metabolites are almost
     devoid of any estrogenic property, they may be useful for
     long-term anti-angiogenetic therapy in both men and women.
     should be of interest to clin. pharmacol. research since it points to
     potential new aspects in the treatment of cardiovascular diseases.
     362-07-2, 2-Methoxyestradiol 362-08-3, 2-Methoxyestrone
ΙT
     RL: PAC (Pharmacological activity); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (angiogenesis and anti-angiogenesis effects of
        estradiol and metabolites)
              THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L71
     ANSWER 4 OF 22 HCAPLUS COPYRIGHT 2002 ACS
     2001:360984 HCAPLUS
ΑN
DN
     135:56155
     Oestradiol metabolites: Effects on airway remodelling
TI
     Stewart, A. G.; Vlahos, R.; Fernandes, D. J.; Hughes, R. A.
ΑU
CS
     Department of Pharmacology, University of Melbourne, Australia
     Prog. Respir. Res. (2001), 31 (New Drugs for Asthma, Allergy and COPD),
SO
     102-105
     CODEN: PRRRAE; ISSN: 1422-2140
PB
     S. Karger AG
DT
     Journal; General Review
LA
     English
     A review with 25 refs. Airway wall remodelling is a significant
AB
     contributor to the airway hyperresponsiveness in asthma and hence to
     asthma symptoms. Existing anti-asthma agents (glucocorticoids) appear to
     have insufficient efficacy in the regulation of smooth muscle
```

proliferation and other aspects of airway wall remodelling in severe asthma. Thus, agents that regulate airway wall remodelling will add significantly to the choice of preventative therapies available for more

estradiol metabolite, 2-methoxyestradiol, is a candidate prophylactic

severe asthma. The anti-angiogenic, anti-proliferative

anti-asthma agent that may more specifically target the airway wall remodelling process.

IT 362-07-2, 2-Methoxyestradiol

RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (estradiol metabolites effects on airway remodelling in relation to asthma)

RE.CNT 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L71 ANSWER 5 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:310080 HCAPLUS
- DN 135:116756
- TI The antiangiogenic property of docetaxel is synergistic with a recombinant humanized monoclonal antibody against vascular endothelial growth factor or 2-methoxyestradiol but antagonized by endothelial growth factors
- AU Sweeney, Christopher J.; Miller, Kathy D.; Sissons, Sean E.; Nozaki, Shinichi; Heilman, Douglas K.; Shen, Jianzhao; Sledge, George W., Jr.
- CS Department of Medicine, Indiana University, Indianapolis, IN, 46202, USA
- SO Cancer Res. (2001), 61(8), 3369-3372 CODEN: CNREA8; ISSN: 0008-5472
- PB American Association for Cancer Research
- DT Journal
- LA English
- Numerous chemotherapeutic agents have been shown to have an inhibitory AB effect on endothelial cell proliferation and migration, and tubule In this study, we examd. the antiangiogenic activity formation. of docetaxel. Docetaxel inhibited endothelial cell proliferation and tubule formation in vitro in a dose-dependent fashion. Docetaxel treatment also inhibited angiogenesis in an in vivo Matrigel plug assay. The endothelial stimulating factors, vascular endothelial cell growth factor (VEGF) and basic fibroblast growth factor are able to protect endothelial cells from the antiangiogenic properties of docetaxel. This protective effect can be overcome by a recombinant humanized monoclonal antibody directed against VEGF in both in vitro and in vivo models. Similarly, combination of docetaxel with the antiangiogenic agent 2-methoxyestradiol also overcomes the protective effect of VEGF in both in vitro and in vivo models. suggest that microenvironmental factors (e.g., local release of VEGF and basic fibroblast growth factor) could play a role in decreasing the antiangiogenic effects of docetaxel, whereas agents such as 2-methoxyestradiol and recombinant humanized monoclonal antibody directed against VEGF may reverse this protective effect.
- IT 362-07-2, 2-Methoxyestradiol

RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (antiangiogenic property of docetaxel is synergistic with a humanized MAb against vascular endothelial growth factor or 2-methoxyestradiol but antagonized by endothelial growth factors)

RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L71 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:43056 HCAPLUS
- DN 135:71332
- TI 2-Methoxyestradiol: an endogenous antiangiogenic and antiproliferative drug candidate
- AU Pribluda, Victor S.; Gubish, Edward R., Jr.; LaVallee, Theresa M.; Treston, Anthony; Swartz, Glenn M.; Green, Shawn J.
- CS EntreMed Inc., Rockville, MD, USA
- SO Cancer Metastasis Rev. (2000), 19(1/2), 173-179 CODEN: CMRED4; ISSN: 0167-7659
- PB Kluwer Academic Publishers
- DT Journal; General Review
- LA English

- AB A review, with 43 refs. 2-Methoxyestradiol (2ME2), once considered an inactive end-metabolite of estradiol, has recently emerged as a very promising agent for cancer treatment. It is orally active in a wide range of tumor models, and inhibits tumor growth at doses showing no clin. signs of toxicity. 2ME2 targets both the tumor cell and endothelial cell compartments by inducing apoptosis in rapidly proliferating cells and inhibiting blood vessel formation at several stages in the angiogenic cascade. Moreover, the ability of 2ME2 to inhibit metastatic spread in several models adds to its therapeutic value for cancer treatment at various stages of the disease. Though the mechanism of action is still undefined, several potential mol. targets and pathways of activation have been suggested.
- IT 362-07-2, 2-Methoxyestradiol
 RL: BAC (Biological activity or effector, except adverse); THU
 (Therapeutic use); BIOL (Biological study); USES (Uses)
 (methoxyestradiol as endogenous antiangiogenic and
 antiproliferative drug candidate)
- RE.CNT 43 THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L71 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- AN 2000:595071 HCAPLUS
- DN 133:305736
- TI 2-Methoxyestradiol blocks **estrogen**-induced rat pituitary tumor growth and tumor **angiogenesis**: possible role of vascular endothelial growth factor
- AU Banerjee, Sushanta K.; Zoubine, Mikhail N.; Sarkar, Dipak K.; Weston, Allan P.; Shah, Jamshed H.; Campbell, Donald R.
- CS Department of Medicine, University of Kansas Medical Center, Kansas City, KS, USA
- SO Anticancer Res. (2000), 20(4), 2641-2645 CODEN: ANTRD4; ISSN: 0250-7005
- PB International Institute of Anticancer Research
- DT Journal
- LA English
- Natural and synthetic estrogens have been assocd. with several types of human and animal cancers including prolactin-secreting pituitary tumors in Fischer 344 rats. These prolactin-secreting tumors are highly angiogenic and their growth is angiogenic dependent. In the present study we have utilized this model to evaluate the effect of 2-methoxyestradiol (2-ME), an endogenous estrogen metabolite that is a potent inhibitor of endothelial cell proliferation in vitro, on estrogen-induced pituitary tumor growth and angiogenesis
 - Adult female rats were implanted (s.c.) with a silastic capsule contg. estradiol-17.beta. (E2). After seven days of const. E2 exposure animals were injected (s.c.) daily with 25 mg/kg of 2-ME and killed either three or 8 days later. Changes in pituitary wt. and proliferating cell nuclear antigen (PCNA) labeling index indicated growth while degree of angiogenesis was detd. immunohistochem. using factor VIII related The results indicate that 2-ME inhibited estrogen antigen. -induced lactotroph growth by 32% and tumor angiogenesis by 89%. Furthermore, vascular endothelial growth factor (VEGF) expression, evaluated by immunohistochem. anal., was down-regulated concomitant with tumor angiogenic suppression. These studies suggest that 2-ME may have therapeutic potential for hormone-induced cancer and that its angiostatic activity may be modulated through down-regulation of VEGF expression.
- IT 362-07-2, 2-Methoxyestradiol
 - RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (methoxyestradiol blocks estrogen-induced rat pituitary tumor
- growth and tumor **angiogenesis** and VEGF role therein)
 RE.CNT 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

```
2000:592560 HCAPLUS
AN
DN
     133:198575
     Compositions and methods for use in targeting vascular destruction
ΤI
     Pero, Ronald W.; Sherris, David
ΙN
PΑ
     Oxigene, Inc., USA
     PCT Int. Appl., 36 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                     KIND DATE
                                           APPLICATION NO.
                                                            DATE
     PATENT NO.
                                           _____
     _____
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                           _____
                                                            20000216
                            20000824
                                           WO 2000-US3996
     WO 2000048606
PΙ
                      Α1
        W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN,
             IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
            MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
             SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                            20000216
                                          EP 2000-914606
     EP 1152764
                       Α1
                            20011114
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
PRAI US 1999-120478
                      Ρ
                            19990218
                            20000216
     WO 2000-US3996
                       W
OS
     MARPAT 133:198575
     Treatment of warm-blooded animals having a tumor or non-malignant
AB
     hypervascularization, by administering a sufficient amt. of a cytotoxic
     agent formulated into a phosphate prodrug form having substrate
     specificity for microvessel phosphatases, so that microvessels are
     destroyed preferentially over other normal tissues, because the less
     cytotoxic prodrug form is converted to the highly cytotoxic
     dephosphorylated form.
     362-07-2, 2-Methoxyestradiol
ΙT
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (prodrugs for use in targeting vascular destruction)
              THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 9 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
     2000:261865 HCAPLUS
AN
DN
TI
     Enhancement of radiation effects in vitro by the estrogen
     metabolite 2-methoxyestradiol
ΑU
     Amorino, George P.; Freeman, Michael L.; Choy, Hak
     Department of Radiation Oncology, Vanderbilt University Medical Center,
CS
     Nashville, TN, 37232, USA
     Radiat. Res. (2000), 153(4), 384-391
SO
     CODEN: RAREAE; ISSN: 0033-7587
PB
     Radiation Research Society
DT
     Journal
LA
     2-Methoxyestradiol (2-ME) is an endogenous estradiol metabolite that
AB
     disrupts microtubule function, suppresses murine tumors, and inhibits
                   Since some microtubule inhibitors have been shown
     to alter radiosensitivity, we have evaluated 2-ME as a radiation enhancer
     in vitro. H460 human lung cancer cells were plated, treated with 2-ME for
     24 h, and irradiated; then colony-forming ability was assessed.
     radiation dose enhancement ratios (DERs) using this protocol were 1.3, 1.8
     and 2.1 for 1, 1.5 and 2 .mu.M 2-ME, resp. Using a single-cell plating
     protocol, the resp. DERs were 1.2, 1.5 and 1.8. The parent compd. of
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2-ME, .beta.-estradiol, did not enhance radiation effects at equally

cytotoxic doses. Isobologram anal. showed that 1 .mu.M 2-ME was additive with radiation, but that 1.5 and 2 .mu.M were synergistic. Cell cycle anal. showed a dose-dependent increase in the percentage of cells in the radiosensitive G2/M phase after a 24-h treatment with 2-ME; a threefold increase in the percentage of cells in G2/M phase was obsd. using 2 .mu.M Treatment with 2 .mu.M 2-ME almost completely inhibited repair of sublethal damage (SLD) as shown using split-dose recovery. Radiosensitive, repair-deficient murine SCID (severe combined immunodeficient) cells did not show enhancement of radiation effects with 2 .mu.M 2-ME, but enhancement was obsd. in the wild-type parental cells (CB-17). SCID cells complemented with human DNA-dependent protein kinase restored radioenhancement by 2-ME. In addn., MCF-7 breast cancer cells were also radiosensitized by 2 .mu.M 2-ME (DER = 2.1). These data suggest that 2-ME is a potential radiation sensitizer, in addn. to its previously reported antitumor and antiangiogenic properties. We have verified the antiangiogenic activity of 2-ME in vitro using human endothelial cells. Based on these results, we hypothesize that the mechanism of radiation enhancement may involve redistribution of cells into G2/M phase by 2-ME, and that the resulting population of cells is repair-deficient and thus radiosensitive. 362-07-2, 2-Methoxyestradiol RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (estrogen metabolite 2-methoxyestradiol enhancement of radiation effects) THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 36 ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 10 OF 22 HCAPLUS COPYRIGHT 2002 ACS 2000:227537 HCAPLUS 132:262172 Use of neoangiogenesis markers for diagnosis and treatment of Krause, Werner; Muschick, Peter Schering Aktiengesellschaft, Germany PCT Int. Appl., 27 pp. CODEN: PIXXD2 Patent German FAN.CNT 1 KIND DATE APPLICATION NO. DATE PATENT NO. ____ WO 2000018439 A2 20000406 WO 1999-EP7198 19990929 20000914 WO 2000018439 А3 AE, AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CR, CU, CZ, DM, EE, ES, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE 20000413 DE 1998-19845798 19980929 DE 19845798 A1 PRAI DE 1998-19845798 A 19980929 Neoangiogenesis markers (i.e. antibodies or receptors for e.g. vascular endothelial growth factor, placenta growth factor, acidic or basic FGF, transforming growth factor .alpha. or .beta., hepatocyte growth factor, insulin-like growth factor I, glycoprotein B61, protein LERK-1, flk-1 receptor, etc.) or partial sequences thereof and antiangiogenic compds. and factors such as paclitaxel, endostatin, fibronectin peptide, and fumagillin are conjugated with active agents such as chemotherapeutic agents, radiosensitizers, photosensitizers, antibodies, oligonucleotides, radioactive metal complexes, etc., which may be bound to carriers, for treatment of tumors. Likewise, neoangiogenesis markers may be conjugated to diagnostic agents such as MRI, radiog., ultrasound, or near-IR contrast agents for tumor

diagnosis. Thus, N', N', N''', N'''-tetrakis(tert-butoxycarboxymethyl)-N''-

IT

L71 AN

DN

ΤI

ΙN

PA

SO

DT

LA

ΡI

AΒ

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(hydroxycarboxymethyl)diethylenetriamine was converted to its
     N-hydroxysuccinimide ester, coupled to a Thy-1 antibody, complexed with
     186Re, and injected i.v. into rabbits for detection of implanted VX2
     tumors by scintigraphy with a gamma camera.
ΙT
     362-07-2
    RL: ARG (Analytical reagent use); BAC (Biological activity or effector,
     except adverse); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (angiogenesis marker; use of neoangiogenesis
       markers for diagnosis and treatment of tumors)
    ANSWER 11 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1.71
     2000:144722 HCAPLUS
ΑN
     132:185454
DN
     Use of anti-angiogenic agents for inhibiting vessel wall injury
ΤI
     Brown, Charles L., III; Gorlin, Steve
ΙN
     Global Vascular Concepts, Inc., USA
PΑ
     PCT Int. Appl., 29 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                    KIND DATE
     PATENT NO.
                                         APPLICATION NO. DATE
                    ____
                                          WO 1999-US19218 19990824
    WO 2000010552
                      A2
                           20000302
PΙ
                     A3
                           20001123
    WO 2000010552
        W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
            CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
             IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD,
            MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
             SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG,
             KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
             ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
            CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     AU 9956871
                           20000314
                                         AU 1999-56871
                                                           19990824
                     A1
                           19980824
PRAI US 1998-97579
                      Ρ
                           19990824
    WO 1999-US19218
                      W
     Use of anti-angiogenic agents to inhibit an undesirable response
AΒ
     to vessel wall injury, including stent neointima, dialysis graft
     neointima, vascular graft-induced neointima, and the treatment of benign
     hypertrophic scar formation as well as the treatment and passivation of
     unstable atherosclerotic plaques are provided. The invention provides for
     the use of catheter-based devices for enhancing the local delivery of
     anti-angiogenic agents into the endothelial tissues of blood
     vessels of the living body.
IT
     362-07-2, 2-Methoxyestradiol
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (anti-angiogenic agents for inhibiting vessel wall injury)
    ANSWER 12 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
     1999:736476 HCAPLUS
AN
     131:346535
DN
ΤI
     Use of neomycin for treating angiogenesis-related diseases
     Hu, Guo-Fu; Vallee, Bert L.
IN
     The Endowment for Research In Human Biology, Inc., USA
PA
     PCT Int. Appl., 74 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                     KIND DATE
                                          APPLICATION NO. DATE
     PATENT NO.
                                          -----
                           _____
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                                          WO 1999-US10269 19990511
                            19991118
ΡI
     WO 9958126
                      A1
         W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
             DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
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JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
             MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,
             MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
             ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
             CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                          AU 1999-39804
                                                            19990511
     AU 9939804
                      Α1
                            19991129
                                           EP 1999-922915
                                                            19990511
                            20010321
     EP 1083896
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
         R:
             IE, FI
                            19980511
PRAI US 1998-84921
     WO 1999-US10269
                       W
                            19990511
     The present invention is directed to using neomycin or an analog thereof
AB
     as a therapeutic agent to treat angiogenesis-related diseases,
     which are characterized by excessive, undesired or inappropriate
     angiogenesis or proliferation of endothelial cells. The present
     invention is also directed to pharmaceutical compns. comprising: (a)
     neomycin or an analog and, optionally, (b) another anti-angiogenic
     agent or an anti-neoplastic agent. The present invention is further
     directed to a method for screening neomycin analogs having anti-
     angiogenic activity. A preferred embodiment of the invention
     relates to using neomycin to treat subjects having such diseases. A dose
     of 20 ng neomycin/embryo or higher completely inhibited angiogenin
     -induced angiogenesis in the chorioallantoic membrane (CAM)
     assay. Neomycin inhibits angiogenin-induced
     angiogenesis mainly through inhibition of nuclear translocation of
     angiogenin.
     362-07-2, 2-Methoxyestradiol
ΙT
     RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (neomycin, its analogs and other agents for treatment of
        angiogenesis-related diseases)
              THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 13 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
     1999:672339 HCAPLUS
ΑN
     132:520
DN
     Angiostatic activity of steroids in the chick embryo CAM and
TI
     rabbit cornea models of neovascularization
ΑU
     McNatt, Loretta G.; Weimer, Lori; Yanni, John; Clark, Abbot F.
     Alcon Laboratories, Inc., Fort Worth, TX, USA
CS
     J. Ocul. Pharmacol. Ther. (1999), 15(5), 413-423
SO
     CODEN: JOPTFU; ISSN: 1080-7683
PΒ
     Mary Ann Liebert, Inc.
DΤ
     Journal
LA
     English
     Ocular neovascular diseases represent a major cause of blindness in the
AΒ
            Angiostatic steroids are a unique class of compds. which
     inhibit the formation of new blood vessels in various models, including
     ocular models of angiogenesis. In search of potent new anti-
     angiogenic agents for the treatment of ocular neovascular disease,
     a large group of steroids were evaluated for angiostatic
     activity in the chick embryo CAM model. Angiostatic activity
     was found among all steroid classes included in the study.
                                                                 There was a
     good correlation between the angiostatic efficacies of 15
     diverse steroids tested in the chick CAM and in the rabbit LPS-induced
     corneal pocket models of neovascularization (r=0.76, p=0.01). These
     studies show that potent angiostatic steroids inhibit
     neovascularization in two different animal models, suggesting a common
     mechanism of action. Glucocorticoid therapy is sometimes assocd. with
     ocular side effects. Two of the most potent angiostatic
     steroids, AL-3789 and AL-4940, were evaluated for glucocorticoid-mediated
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anti-inflammatory activity in the in vitro U937 cell model of LPS-induced

IL-1 induction and found to be devoid of glucocorticoid activity.

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Angiostatic steroids which lack glucocorticoid activity should be
     attractive drug candidates for treating ocular neovascular disease.
ΙT
     362-07-2, 2-Methoxyestradiol
     RL: BAC (Biological activity or effector, except adverse); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study);
     USES (Uses)
        (angiostatic activity of steroids in chick embryo CAM and
        rabbit cornea models of neovascularization)
              THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
AN
     1999:48631 HCAPLUS
DN
     130:119599
     Pharmaceutical compositions comprising an angiostatic steroid
ΤI
     combined with a hyaluronan for increasing neovascularization and
     angiogenesis during wound healing
     Seed, Michael P.; Alam, Chandan; Willoughby, Derek A.
IN
PA
     Hyal Pharmaceutical Corporation, Can.
SO
     PCT Int. Appl., 84 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 1
                                           APPLICATION NO.
                                                           DATE
     PATENT NO.
                     KIND DATE
                                           _____
                                                            _____
                                           WO 1998-CA649
                                                            19980703
     WO 9901142
                      A1
                            19990114
PΙ
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
             CM, GA, GN, ML, MR, NE, SN, TD, TG
                                           CA 1997-2208916 19970703
     CA 2208916
                            19990103
                       AA
     AU 9882021
                                           AU 1998-82021
                                                            19980703
                       A1
                            19990125
PRAI CA 1997-2208916
                            19970703
                            19980703
     WO 1998-CA649
     A pharmaceutical compn. is disclosed for increasing neovascularization and
AB
     angiogenesis during wound healing in a mammal beyond the level of
     neovascularization and angiogenesis which would occur at the
     wound site without any treatment, the compn. comprising an effective amt.
     of any angiostatic steroid which has reduced or no deteriorative
     or detrimental side effects, combined with an effective amt. of a form of
     hyaluronan, e.g. hyaluronic acid or a pharmaceutically acceptable salt
     thereof.
ΙT
     362-07-2 362-08-3
     RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (angiostatic steroid-hyaluronan combination for increasing
        neovascularization and angiogenesis during wound healing)
              THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 15 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
     1998:785662 HCAPLUS
ΑN
DN
     130:33040
     Methods using 7-(substituted amino)-9-[(substituted glycyl)amido]-6-
ΤI
     demethyl-6-deoxytetracyclines for inhibiting angiogenesis,
     proliferation of endothelial or tumor cells, and tumor growth
     Backer, Joseph M.; Bohlen, Peter
IN
     American Cyanamid Company, USA
PA
SO
     U.S., 12 pp.
     CODEN: USXXAM
```

DT

Patent

```
gazi - 09 / 899702
LA
    English
FAN.CNT 1
                                   APPLICATION NO. DATE
                  KIND DATE
    PATENT NO.
                     ____
    US 5843925 A 19981201
US 5856315 A 19990105
                                         US 1994-354694
                                                           19941213
PΙ
                                        US 1998-84484 19980526
PRAI US 1994-354694
                          19941213
    MARPAT 130:33040
    A method is provided for inhibiting angiogenesis and
AΒ
    proliferation of endothelial cells by administering an inhibitory amt. of
    a 7-(substituted amino)-9-[(substituted glycyl)amido]-6-demethyl-6-
   . deoxytetracycline (Markush included). Also provided is a method for
     inhibiting proliferation of tumor cells and tumor growth by administering
     an inhibitory amt. of a compd. of the invention in combination with a
     chemotherapeutic agent or radiation therapy. Further provided are compns.
     contg. an effective inhibitory amt. of a compd. of the invention in a
    pharmaceutically acceptable carrier.
    362-07-2, 2-Methoxyestradiol
    RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (deoxytetracycline prepn. for inhibiting angiogenesis,
       proliferation of endothelial or tumor cells, and tumor growth, and
       compns. with other agents)
             THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
```

L71 ANSWER 16 OF 22 HCAPLUS COPYRIGHT 2002 ACS

AN 1998:778968 HCAPLUS

DN 130:105451

TI Inhibition of normal and experimental **angiotumor** endothelial cell proliferation and cell cycle progression by 2-methoxyestradiol

AU Reiser, F.; Way, D.; Bernas, M.; Witte, M.; Witte, C.

CS Department of Surgery, The University of Arizona, Tucson, AZ, 85724, USA

SO Proc. Soc. Exp. Biol. Med. (1998), 219(3), 211-216 CODEN: PSEBAA; ISSN: 0037-9727

PB Blackwell Science, Inc.

DT Journal

LA English

With rapid growth and metab., aggressive cancers require an extensive . AB vascular network, termed tumor angiogenesis. The body produces a variety of natural angiogenic inhibitors, among which is the mammalian estrogen metabolite, 2-methoxyestradiol (2-MeOE2). In this study, we compared the effects of 2-MeOE2 on a human umbilical vein cell line (HUVEC-C) and on an immortal, angiotumor-producing rat sinusoidal endothelial cell line (RSE-1). In vitro, the effects of varying concns. of 2-MeOE2 from 0.01-100.0 .mu.M were measured with cell counts and compared to control cells. HUVEC-C had an ED50 .apprx.3.5 .mu.M with .apprx.27% inhibition of cell growth whereas RSE-1 had an ED50 .apprx.2.2 .mu.M with .apprx.50% inhibition of cell growth compared with controls. The lowest concn. with maximal effect was 10.0 .mu.M 2-MeOE2 for both cell lines. Using this concn., flow cytometric anal. of cell cycles was performed with propidium iodide stained DNA of HUVEC-C and RSE-1 at 24 and 48 h. Both demonstrated a significant block at G2M of the cell cycle. At 48 h, HUVEC-C had 32% of cells in G2M (control = 9% G2M), and RSE-1 had 36% of cells in G2M (control = 18% G2M). These findings demonstrate a strong in vitro antiproliferative effect of 2-MeOE2 on normal dividing endothelial as well as angiotumor cells mediated through a cell cycle-specific block at G2M. The antiendothelial, antiangiotumor effect of 2-MeOE2 supports its potential as a therapeutic agent against solid organ cancers, benign or malignant vascular growths, and other pathol. states dependent on angiogenesis.

IT 362-07-2, 2-Methoxyestradiol
RL: BAC (Biological activity or effector, except adverse); THU
(Therapeutic use); BIOL (Biological study); USES (Uses)
(methoxyestradiol inhibition of normal and exptl. angiotumor

endothelial cell proliferation and cell cycle progression)
RE.CNT 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

```
ANSWER 17 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
    1998:369884 HCAPLUS
ΑN
DN
     129:90490
     Is 2-methoxyestradiol an endogenous estrogen metabolite that
ΤI
     inhibits mammary carcinogenesis?
ΑU
     Zhu, Bao Ting; Conney, Allan H.
     Laboratory for Cancer Research, Department of Chemical Biology, College of
CS
     Pharmacy, Rutgers-The State University of New Jersey, Piscataway, NJ,
     08854-8020, USA
     Cancer Res. (1998), 58(11), 2269-2277
SO
     CODEN: CNREA8; ISSN: 0008-5472
    American Association for Cancer Research
PB ·
     Journal; General Review
DΤ
LA
    English
    A review, with 172 refs. Catechol estrogens (2- or
AB
     4-hydroxyestradiol and 2- or 4-hydroxyestrone) are chem. reactive
     estrogen metabolites that are O-methylated to less polar
    monomethyl ethers by catechol-O-methyltransferase, an enzyme present in
    many tissues such as the liver, kidney, brain, placenta, uterus, and
    mammary gland. In the present report, the authors review recent studies
     on the antitumorigenic and antiangiogenic effects of exogenously
     administered 2-methoxyestradiol in vitro and in vivo. The authors also
     discuss data that suggest that endogenous formation of 2-methoxyestradiol
     (and its 2-hydroxyestradiol precursor) may have a protective effect on
     estrogen-induced cancers in target organs. Although the mol.
     mechanism of action of 2-methoxyestradiol is not clear, the authors
     suggest that some unique effects of 2-methoxyestradiol may be mediated by
     a specific intracellular effector or receptor that is refractory to the
     parent hormone, estradiol. Addnl. research is needed to identify factors
     that regulate the metabolic formation and disposition of
     2-methoxyestradiol in liver and in target cells and to evaluate the
     effects of modulating 2-methoxyestradiol formation on estrogen
     -induced carcinogenesis.
ΙT
     362-07-2, 2-Methoxyestradiol
     RL: BAC (Biological activity or effector, except adverse); MFM (Metabolic
     formation); THU (Therapeutic use); BIOL (Biological study); FORM
     (Formation, nonpreparative); USES (Uses)
        (methoxyestradiol antitumorigenic and antiangiogenic effects
        in vitro and in vivo)
    ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L71
AN
     1997:752842 HCAPLUS
DN
     128:30711
     Treatment of asthma and airway diseases using steroids or steroid analogs
ΤI
IN
     Stewart, Alastair George
     Amrad Operations Pty. Ltd., Australia; Stewart, Alastair George
PA
SO
     PCT Int. Appl., 63 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
                                         APPLICATION NO. DATE
                     KIND DATE
     PATENT NO.
     ______
                                          _____
                      A1
                           19971120
                                         WO 1997-AU286
                                                          19970509
PΙ
     WO 9742958
            AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
             DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ,
             LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL,
             PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ,
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RW: GH, KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

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19971205
      AU 9726276
                      A1
                                            AU 1997-26276
                                                             19970509
      AU 725048
                        В2
                             20001005
                                            CN 1997-194505
                                                             19970509
                        Α
                             19990602
      CN 1218401
                        A1
                                            EP 1997-917945
                             19990623
                                                             19970509
      EP 923376
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, FI
                                            US 1997-853528
      US 5962445
                                                             19970509
                             19991005
                        Α
      JP 2000507923
                       Т2
                             20000627
                                            JP 1997-530434
                                                             19970509
                        В1
                             20010313
                                            US 1999-330918
                                                             19990611
      US 6200966
                        Α
 PRAI AU 1996-9766
                             19960509
                        Α
                             19960520
      AU 1996-9918
      US 1997-853528
                        A1
                             19970509
                      W
                             19970509
      WO 1997-AU286
      The invention relates to use of steroids or steroid analogs in the
· AB
      treatment of chronic and acute inflammation of the airways, particularly
      asthmatic conditions. It also relates to compds. and compns. which
      modulate airway remodelling. In a preferred embodiment, the active
      component inhibits inflammation and smooth muscle cell proliferation in
      the airway wall. It may also have a least one other activity selected
      from antiangiogenesis, antioxidn., and the ability to disrupt
      microtubule formation. In a preferred embodiment, the steroid is
      2-methoxyoestradiol.
      362-07-2, 2-Methoxyestradiol 362-08-3, 2-Methoxyestrone
 ΙT
      RL: BAC (Biological activity or effector, except adverse); THU
      (Therapeutic use); BIOL (Biological study); USES (Uses)
         (treatment of asthma and airway diseases using steroids or steroid
         analogs)
      ANSWER 19 OF 22 HCAPLUS COPYRIGHT 2002 ACS
 L71
      1997:483454 HCAPLUS
 AN
      127:91017
 DN
      Method and composition for treatment of pathological conditions associated
 ΤI
      with angiogenesis
      Fotsis, Theodore; Adlercreutz, Herman; Schweigerer, Lothar
 IN
 PA
      Germany
      U.S., 10 pp. Cont.-in-part of U.S. Ser. No. 84,969, abandoned.
 SO
      CODEN: USXXAM
 DT
      Patent
 LA
      English
 FAN.CNT 1
                      KIND DATE
                                            APPLICATION NO. DATE
      PATENT NO.
                            -----
      _____
                       ----
      US 5643900
                        Α
                             19970701
                                            US 1995-405776
                                                             19950317
 PΙ
 PRAI US 1993-84969
                             19930702
      The present invention relates to a method and a compn. for the treatment
      of pathol. conditions assocd. with enhanced angiogenesis. The
      method comprises administering 2-methoxyestradiol to a subject in need of
      such treatment. 2-Methoxyestradiol also has potent pharmacol. properties
      in the treatment of solid tumors.
 TT
      362-07-2, 2-Methoxyestradiol
      RL: BAC (Biological activity or effector, except adverse); THU
      (Therapeutic use); BIOL (Biological study); USES (Uses)
         (treatment of angiogenic diseases and solid tumors with
         2-methoxyestradiol)
      ANSWER 20 OF 22 HCAPLUS COPYRIGHT 2002 ACS
 L71
 AN
      1997:84466 HCAPLUS
 DN
      126:140011
      Suppression of type II collagen-induced arthritis by the endogenous
 TТ
      estrogen metabolite 2-methoxyestradiol
 ΑU
      Josefsson, Elisabet; Tarkowski, Andrej
 CS
      University of Goteborg, Goteborg, Swed.
 SO
      Arthritis Rheum. (1997), 40(1), 154-163
      CODEN: ARHEAW; ISSN: 0004-3591
```

PB

DT

Lippincott-Raven

Journal

- LA English
- To evaluate the antiarthritic properties of 2-methoxyestradiol, an AB endogenous metabolite of estradiol, on type II collagen-induced arthritis (CIA) in DBA/1 mice, the effects of treatment with 2-methoxyestradiol on the development of CIA were evaluated clin. and histol. The in vitro effects of 2-methoxyestradiol on lymphocyte and endothelial cell proliferation and differentiation were analyzed by std. methods. development of CIA was significantly suppressed by 2-methoxyestradiol. Incubation with 2-methoxyestradiol suppressed the in vitro proliferation of endothelial cells, indicating that this compd. down-regulates angiogenesis. Endothelial cell prodn. of nitric oxide (NO) was also down-regulated by 2-methoxyestradiol. In contrast to estradiol, 2-methoxyestradiol exerted neither detectable feminizing effects on the sex organs nor inhibition of leukocyte development in hematopoietic organs. The development of CIA is suppressed by 2-methoxyestradiol, possibly via inhibition of angiogenesis. Diminished NO prodn. could be of importance in vivo because it is a potent pro-inflammatory mediator. Since 2-methoxyestradiol exerts only mild side effects compared with estradiol, it is an interesting candidate for therapeutic use in inflammatory diseases.
- IT 362-07-2, 2-Methoxyestradiol
 - RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (suppression of type II collagen-induced arthritis by endogenous estrogen metabolite 2-methoxyestradiol)
- L71 ANSWER 21 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- AN 1997:39755 HCAPLUS
- DN 126:127078
- TI Inhibition of **angiogenesis** and breast cancer in mice by the microtubule inhibitors 2-methoxyestradiol and taxol
- AU Klauber, Nancy; Parangi, Sareh; Flynn, Evelyn; Hamel, Ernest; D'Amato, Robert J.
- CS Department Surgery, Children's Hospital and Harvard Medical School, Boston, MA, 02115, USA
- SO Cancer Res. (1997), 57(1), 81-86 CODEN: CNREA8; ISSN: 0008-5472
- PB American Association for Cancer Research
- DT Journal
- LA English
- 2-Methoxyestradiol (2-ME), an endogenous estrogen metabolite AB which disrupts microtubule function, has been shown to inhibit proliferating cells in vitro and suppress certain murine tumors in vivo. In vitro screening has detd. that breast cancer cell lines are most sensitive to inhibition by 2-ME. Addnl., 2-ME has been shown to inhibit angiogenesis in vitro. We tested whether 2-ME suppresses cytokine-induced angiogenesis in vivo and inhibits growth of a human breast carcinoma in severe combined immunodeficient mice. A model of basic fibroblast growth factor (bFGF) and vascular endothelial growth factor (VEGF)-induced corneal neovascularization in C57BL/6 mice was used to evaluate the antiangiogenic effects of 2-ME and other microtubule inhibitors such as Taxol, vincristine, and colchicine. 2-ME (150 mg/kg p.o.) inhibited bFGF and VEGF-induced neovascularization by 39% and 54%, resp. Taxol (6 mg/kg i.p.) inhibited bFGF and VEGF-induced neovascularization by 45% and 37%, resp. Vincristine (0.2 mg/kg i.p.) and colchicine (0.25 mg/kg i.p.) had no effect. Treatment with 2-ME (75 mg/kg p.o.) for 1 mo suppressed the growth of a human breast carcinoma in mice by 60% without toxicity. Recognition of the antiangiogenic and antitumor properties of 2-ME and taxol may be crucial in planning clin. applications to angiogenesis-dependent diseases.
- IT 362-07-2, 2-Methoxyestradiol
 - RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (inhibition of angiogenesis and breast cancer in mice by microtubule inhibitors methoxyestradiol and taxol)
- L71 ANSWER 22 OF 22 HCAPLUS COPYRIGHT 2002 ACS

- AN 1996:640895 HCAPLUS
- DN 125:292016
- TI Inhibitors of angiogenesis in human urine
- AU Fotsis, Theodore; Pepper, Michael S.; Aktas, Erkan; Joussen, Antonia; Kruse, Friedrich; Adelcreutz, Herman; Waehaelea, Kristina; Hase, Tapio; Montesano, Roberto; Schweigerer, Lothar
- CS Children's University Hospital, University of Heidelberg, Heidelberg, 69120, Germany
- SO NATO ASI Ser., Ser. A (1996), 285 (Molecular, Cellular, and Clinical Aspects of Angiogenesis), 213-227 CODEN: NALSDJ; ISSN: 0258-1213
- DT Journal; General Review
- LA English
- AB A review with .apprx.50 refs. on the authors' work in screening urine of human subjects consuming a diet rich in plant products for antiangiogenic compds. The authors identified the isoflavonoid genistein, the endogenous estrogen metabolite 2-methoxyestradiol and metabolites of flavonoids that show inhibitory activities against angiogenesis. Irresp. of whether or not these substances play any physiol. role in humans, they might be suitable as pharmacol. agents for treatment of solid malignant tumors and other angiogenic diseases.
- IT 362-07-2, 2-Methoxyestradiol
 RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); THU (Therapeutic use); BIOL (Biological study);
 OCCU (Occurrence); USES (Uses)
 (inhibitors of angiogenesis isolated from human urine in relation to antitumor activity and treatment of angiogenic diseases)

=> fil reg FILE 'REGISTRY' ENTERED AT 15:30:07 ON 19 FEB 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 American Chemical Society (ACS)

STRUCTURE FILE UPDATES: 18 FEB 2002 HIGHEST RN 393508-26-4 DICTIONARY FILE UPDATES: 18 FEB 2002 HIGHEST RN 393508-26-4

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when conducting ${\tt SmartSELECT}$ searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

The P indicator for Preparations was not generated for all of the CAS Registry Numbers that were added to the H/Z/CA/CAplus files between 12/27/01 and 1/23/02. Use of the P indicator in online and SDI searches during this period, either directly appended to a CAS Registry Number or by qualifying an L-number with /P, may have yielded incomplete results. As of 1/23/02, the situation has been resolved. Also, note that searches conducted using the PREP role indicator were not affected.

Customers running searches and/or SDIs in the H/Z/CA/CAplus files incorporating CAS Registry Numbers with the P indicator between 12/27/01 and 1/23/02, are encouraged to re-run these strategies. Contact the CAS Help Desk at 1-800-848-6533 in North America or 1-614-447-3698, worldwide, or send an e-mail to help@cas.org for further assistance or to receive a credit for any duplicate searches.

=> d ide can tot 169

L69 ANSWER 1 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **383414-35-5** REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(2,2,2-trifluoroethyl)-, (17.beta.)-(9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 678473

FS STEREOSEARCH

MF C20 H25 F3 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry.

remaining hil compounds from references

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

L69 ANSWER 2 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **302799-37-7** REGISTRY

CN Estra-1,3,5(10)-triene-3,6,17-triol, 2-ethoxy-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 683688

FS STEREOSEARCH

MF C20 H28 O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 133:322044

L69 ANSWER 3 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 192062-20-7 REGISTRY

CN Estra-1,3,5(10)-trien-6-one, 2-ethoxy-3,17-dihydroxy-, O-methyloxime, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 683125

FS STEREOSEARCH

MF C21 H29 N O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry. Double bond geometry unknown.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)
3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 133:322044

REFERENCE 3: 127:95444

L69 ANSWER 4 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 192062-15-0 REGISTRY

CN Estra-1,3,5(10)-trien-6-one, 3,17-dihydroxy-2-(2,2,2-trifluoroethoxy)-, oxime, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 681683

FS STEREOSEARCH

MF C20 H24 F3 N O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry.

Double bond geometry unknown.

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 133:322044

REFERENCE 3: 127:95444

L69 ANSWER 5 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 192062-14-9 REGISTRY

CN Estra-1, 3, 5(10) -trien-6-one, 2-ethoxy-3, 17-dihydroxy-, oxime, (17.beta.)-

(9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 680185

FS STEREOSEARCH

MF C20 H27 N O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry.

Double bond geometry unknown.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 133:322044

REFERENCE 3: 127:95444

L69 ANSWER 6 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **192062-13-8** REGISTRY

CN Estra-1,3,5(10)-trien-6-one, 3,17-dihydroxy-2-(2,2,2-trifluoroethoxy)-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN NSC 681684

FS STEREOSEARCH

MF C20 H23 F3 O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 133:322044

REFERENCE 3: 127:95444

L69 ANSWER 7 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 192062-12-7 REGISTRY

CN Estra-1,3,5(10)-trien-6-one, 2-ethoxy-3,17-dihydroxy-, (17.beta.)- (9CI)

(CA INDEX NAME)

OTHER NAMES:

CN NSC 679431

FS STEREOSEARCH

MF C20 H26 O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 134:326663

REFERENCE 3: 133:322044

REFERENCE 4: 133:105195

REFERENCE 5: 132:303593

REFERENCE 6: 127:95444

L69 ANSWER 8 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **192062-02-5** REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(1-propynyl)-, (17.beta.)- (9CI) (CA

INDEX NAME)

OTHER NAMES:

CN NSC 682429

FS STEREOSEARCH

MF C21 H26 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry.

$$Me - C = C$$

$$H$$

$$R$$

$$H$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 127:95444

L69 ANSWER 9 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 165619-13-6 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 17-diol, 2-(1-pentenyl)-, [2(E), 17.beta.]- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C23 H32 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.

Double bond geometry as shown.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 123:112496

L69 ANSWER 10 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 165619-12-5 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 17-diol, 2-(1-butenyl)-, [2(E), 17.beta.]- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

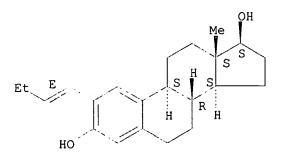
MF C22 H30 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.

Double bond geometry as shown.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 123:112496

L69 ANSWER 11 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **165619-09-0** REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(1-methylethoxy)-, (17.beta.)- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C21 H30 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 123:112496

L69 ANSWER 12 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 165619-08-9 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-propoxy-, (17.beta.)- (9CI) (CA INDEX NAME)

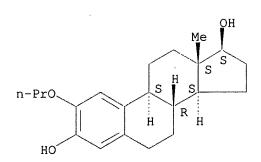
FS STEREOSEARCH

MF C21 H30 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 123:112496

L69 ANSWER 13 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **165619-07-8** REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-ethoxy-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Ethoxyestradiol

CN NSC 671043

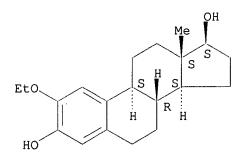
FS STEREOSEARCH

MF C20 H28 O3

SR CA

LC STN Files: BIOSIS, CA, CAPLUS, TOXCENTER, TOXLIT, USPATFULL

Absolute stereochemistry. Rotation (+).



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

7 REFERENCES IN FILE CA (1967 TO DATE)
 7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 134:326663

REFERENCE 3: 133:322044

REFERENCE 4: 132:303593

REFERENCE 5: 129:245330

REFERENCE 6: 127:95444

REFERENCE 7: 123:112496

L69 ANSWER 14 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN **22415-44-7** REGISTRY

CN 19-Norpregna-1,3,5(10)-trien-20-yne-3,17-diol, 2-methoxy-, (17.alpha.)-(9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 19-Nor-17.alpha.-pregna-1,3,5(10)-trien-20-yne-3,17-diol, 2-methoxy- (8CI) OTHER NAMES:

CN 17.alpha.-Ethynyl-2-methoxyestradiol

CN 2-Methoxy-17.alpha.-ethynylestradiol

FS STEREOSEARCH

DR 26011-46-1

MF C21 H26 O3

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT, MEDLINE, TOXCENTER, TOXLIT,

(*File contains numerically searchable property data)

21 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

21 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:64669

REFERENCE 2: 132:132481

REFERENCE 3: 122:256433

REFERENCE 4: 118:161228

REFERENCE 5: 108:94834

REFERENCE 6: 104:1000

REFERENCE 7: 102:106508

REFERENCE 8: 101:48797

REFERENCE 9: 100:185989

REFERENCE 10: 100:17962

L69 ANSWER 15 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 362-08-3 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-methoxy- (8CI, 9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estrone, 2-methoxy- (6CI)

OTHER NAMES:

CN 2-Hydroxyestrone 2-methyl ether

CN 2-Methoxyestrone

FS STEREOSEARCH

MF C19 H24 O3

LC STN Files: ANABSTR, BEILSTEIN*, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CSCHEM, DDFU, DRUGU, EMBASE, MEDLINE, SPECINFO, TOXCENTER, TOXLIT, USPATFULL

(*File contains numerically searchable property data)

Other Sources: EINECS**

(**Enter CHEMLIST File for up-to-date regulatory information)

257 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

257 REFERENCES IN FILE CAPLUS (1967 TO DATE)

34 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:79753

REFERENCE 2: 136:64669

REFERENCE 3: 136:31843

REFERENCE 4: 136:3843

REFERENCE 5: 135:366956

REFERENCE 6: 135:221459

REFERENCE 7: 135:195698

REFERENCE 8: 135:175599

REFERENCE 9: 135:147586

REFERENCE 10: 134:321936

L69 ANSWER 16 OF 16 REGISTRY COPYRIGHT 2002 ACS

RN 362-07-2 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-methoxy-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-triene-3,17.beta.-diol, 2-methoxy- (7CI, 8CI)

CN Estradiol, 2-methoxy- (6CI)

OTHER NAMES:

CN 2-Hydroxyestradiol 2-methyl ether

CN 2-Methoxyestra-1,3,5(10)-triene-3,17.beta.-diol

CN 2-Methoxyestradiol

CN NSC 659853

FS STEREOSEARCH

MF C19 H26 O3

CI COM

LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT,
CBNB, CHEMCATS, CIN, CSCHEM, DDFU, DRUGNL, DRUGU, DRUGUPDATES, EMBASE,
IFICDB, IFIPAT, IFIUDB, MEDLINE, PROMT, RTECS*, SPECINFO, TOXCENTER,
TOXLIT, USPATFULL

(*File contains numerically searchable property data)

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Me OH S S S S R H
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299 REFERENCES IN FILE CA (1967 TO DATE)

7 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

300 REFERENCES IN FILE CAPLUS (1967 TO DATE)

24 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:112738

REFERENCE 2: 136:79991

REFERENCE 3: 136:79754

REFERENCE 4: 136:79753

REFERENCE 5: 136:64669

REFERENCE 6: 136:48639

REFERENCE 7: 136:42803

REFERENCE 8: 136:31870

REFERENCE 9: 136:31843

REFERENCE 10: 136:31835

=> s 130 not 168

L72 64 L30 NOT L68

=>

=> d ide can tot

L72 ANSWER 1 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 345317-45-5 REGISTRY

CN Estra-1,3,5(10)-trien-17-ol, 2-(2-propenyl)-, (17.beta.)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H28 O

SR CA

LC STN Files: CA, CAPLUS, TOXLIT

Absolute stereochemistry.

por structure search

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:56201

L72 ANSWER 2 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 302799-36-6 REGISTRY

CN Estra-1,3,5(10)-trien-3-ol, 17-amino-2-ethoxy-, hydrochloride (9CI) (CA INDEX NAME)

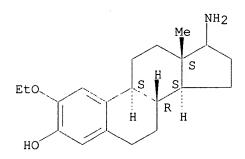
FS STEREOSEARCH

MF C20 H29 N O2 . C1 H

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.



● HCl

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:322044

L72 ANSWER 3 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 300853-53-6 REGISTRY

CN Estra-1, 3, 5(10) - triene-3, 17-diol, 2-methoxy-,

(8.alpha., 9.beta., 13.alpha., 14.beta., 17.alpha.) - (9CI) (CA INDEX NAME)

OTHER NAMES:

CN ent-2-Methoxyestra-1,3,5(10)-triene-16.alpha.,17.beta.-diol

FS STEREOSEARCH

MF C19 H26 O3

SR CA

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:296594

L72 ANSWER 4 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 258278-77-2 REGISTRY

CN Estra-1, 3, 5(10) -triene-2-carbonitrile, 3, 17-dihydroxy-17-phenyl-,

(17.beta.) - (9CI) (CA INDEX NAME)

OTHER NAMES:

CN EM 2318

FS STEREOSEARCH

MF · C25 H27 N O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:152024

L72 ANSWER 5 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 258278-74-9 REGISTRY

CN Estra-1, 3, 5(10) -triene-2-carbonitrile, 3, 17-dihydroxy-17-propyl-,

(17.beta.) - (9CI) (CA INDEX NAME)

OTHER NAMES:

CN EM 2132

FS STEREOSEARCH

MF C22 H29 N O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:152024

L72 ANSWER 6 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 258278-49-8 REGISTRY

CN Estra-1, 3, 5(10) -triene-2-carbonitrile, 3, 17-dihydroxy-17-(phenylmethyl)-,

(17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN EM 1836

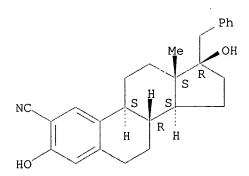
FS STEREOSEARCH

MF C26 H29 N O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 132:152024

L72 ANSWER 7 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 229635-01-2 REGISTRY

CN 19-Norpregna-1,3,5(10),20-tetraene-3,17-diol, 2-methoxy-, (17.alpha.)-

(9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H28 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 135:195698

REFERENCE 2: 131:88084

L72 ANSWER 8 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 192062-30-9 REGISTRY

CN Estra-1,3,5(10)-trien-3-ol, 17-amino-2-ethoxy-, (17.beta.)- (9CI) (CA

INDEX NAME)

FS STEREOSEARCH

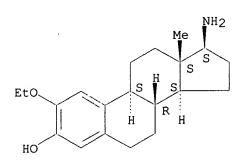
MF C20 H29 N O2

CI COM

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 127:95444

L72 ANSWER 9 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 192062-27-4 REGISTRY

CN Estra-1,3,5(10)-trien-3-ol, 17-amino-2-ethoxy-, hydrochloride, (17.beta.)-(9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H29 N O2 . C1 H

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

CRN (192062-30-9)

Absolute stereochemistry.

HC1

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 127:95444

L72 ANSWER 10 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 192062-18-3 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 6, 17-triol, 2-ethoxy-, (6.alpha., 17.beta.) - (9CI)

(CA INDEX NAME)

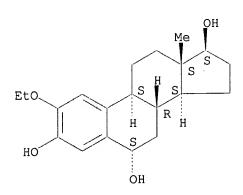
FS STEREOSEARCH

MF C20 H28 O4

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:322044

REFERENCE 2: 127:95444

L72 ANSWER 11 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 192062-17-2 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 6-amino-2-ethoxy-, (17.beta.)- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C20 H29 N O3

SR CA LC STN Files: CA, CAPLUS, TOXCENTER Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 127:95444

L72 ANSWER 12 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 192062-07-0 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(2,2,2-trifluoroethoxy)-, (17.beta.)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H25 F3 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 133:322044

REFERENCE 2: 129:245330

REFERENCE 3: 127:95444

L72 ANSWER 13 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 161762-25-0 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(3-aminopropyl)-, (17.beta.)- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C21 H31 N O2

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

$$H_2N$$
 (CH₂)₃ H_2 H_3 H_4 H_5 H_6 H_7 H_8 H_8

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 122:187857

L72 ANSWER 14 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 161762-24-9 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 17-diol, 2-(3-hydroxypropyl)-, (17.beta.)- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C21 H30 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 136:31870

REFERENCE 2: 124:289981

REFERENCE 3: 122:187857

L72 ANSWER 15 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 161762-23-8 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(2-aminoethyl)-, (17.beta.)- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH

MF C20 H29 N O2

SR CA

CA, CAPLUS, TOXCENTER, TOXLIT LCSTN Files:

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 125:105405

122:187857 REFERENCE 2:

L72 ANSWER 16 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN161762-22-7 REGISTRY

Estra-1,3,5(10)-triene-3,17-diol, 2-(2-hydroxyethyl)-, (17.beta.)- (9CI) CN

(CA INDEX NAME)

FS STEREOSEARCH

MF C20 H28 O3

SR CA

CA, CAPLUS, TOXCENTER, TOXLIT LC STN Files:

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 136:31870 REFERENCE

124:289981 REFERENCE 2:

REFERENCE 3: 122:187857

ANSWER 17 OF 64 REGISTRY COPYRIGHT 2002 ACS L72

RN161762-21-6 REGISTRY

Estra-1,3,5(10)-triene-3,17-diol, 2-(aminomethyl)-, hydrochloride, CN

(17.beta.) - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H27 N O2 . Cl H

SR CA

CA, CAPLUS STN Files: LC

Absolute stereochemistry.

HCl

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 122:187857

ANSWER 18 OF 64 REGISTRY COPYRIGHT 2002 ACS L72

156669-41-9 REGISTRY RN

19-Norpregna-1,3,5(10)-trien-20-yne-3,17-diol, 4-bromo-2-methoxy-, CN

(17.alpha.) - (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Methoxy-4-bromo-17.alpha.-ethynylestradiol

4-Bromo-2-methoxyethinylestradiol CN

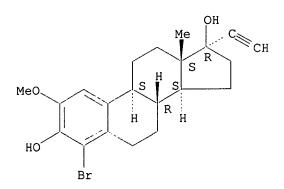
FS STEREOSEARCH

C21 H25 Br O3 MF

SR

LC STN Files: CA, CAPLUS, TOXLIT

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1: 121:100057 REFERENCE

ANSWER 19 OF 64 REGISTRY COPYRIGHT 2002 ACS L72

RN 153654-78-5 REGISTRY CN 19-Norpregna-1,3,5(10)-trien-20-yne-3,17-diol, 2-(hydroxymethyl)-, (17.alpha.)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H26 O3

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: .120:192069

L72 ANSWER 20 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 144216-19-3 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2,4-bis(hydroxymethyl)- (9CI) (CA INDEX NAME)

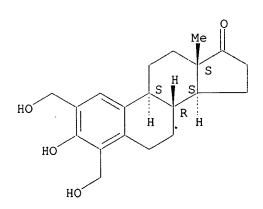
FS STEREOSEARCH

MF C20 H26 O4

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 117:212773

L72 ANSWER 21 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 144216-18-2 REGISTRY

CN 19-Norpregna-1, 3, 5(10) -trien-20-yne-2, 4-dimethanol, 3, 17-dihydroxy-,

(17.alpha.) - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C22 H28 O4

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 120:192069

REFERENCE 2: 117:212773

L72 ANSWER 22 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 144216-17-1 REGISTRY

CN Estra-1, 3, 5(10) -triene-2, 4-dimethanol, 3, 17-dihydroxy-, (17.beta.) - (9CI)

(CA INDEX NAME)

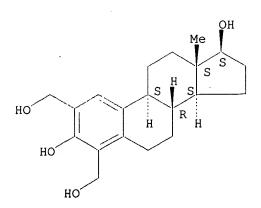
FS STEREOSEARCH

MF C20 H28 O4

SR CA

LC STN Files: CA, CAPLUS, CASREACT

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 120:192069

REFERENCE 2: 117:212773

L72 ANSWER 23 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 141056-21-5 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 4-fluoro-3-hydroxy-2-methoxy- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Methoxy-4-fluoroestrone

CN 4-Fluoro-2-methoxyestrone

FS STEREOSEARCH

MF C19 H23 F O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 116:207984

L72 ANSWER 24 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 119829-59-3 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 17-diol, 2-(1-hydroxy-1-methylethyl)-,

(17.beta.) - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H30 O3

SR CA

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT

(*File contains numerically searchable property data)

1 REFERENCES IN FILE CA (1967 TO DATE)
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 110:154665

L72 ANSWER 25 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 111443-37-9 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-(hydroxymethyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estrone, 2-(hydroxymethyl)- (6CI)

FS STEREOSEARCH

MF C19 H24 O3

SR CAOLD

LC STN Files: CA, CAOLD, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 120:192069

L72 ANSWER 26 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 98543-85-2 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-(1-methyl-2-propenyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-(1-methylallyl)- (7CI)

FS STEREOSEARCH

DR 175414-64-9

MF C22 H28 O2

SR CAOLD

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS

(*File contains numerically searchable property data)

$$H_2C$$
 H_2C
 H_2C
 H_3
 H_4
 H_5
 H_6
 H_7
 H_8
 H

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 124:260811

L72 ANSWER 27 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 97515-50-9 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 4-bromo-2-methoxy-, (17.beta.)- (9CI)

(CA INDEX NAME)

OTHER NAMES:

CN 4-Bromo-2-methoxyestradiol

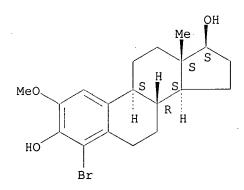
FS STEREOSEARCH

MF C19 H25 Br O3

SR CA

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT, TOXCENTER, TOXLIT (*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 111:233338

REFERENCE 2: 103:65176

L72 ANSWER 28 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 97515-47-4 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 4-fluoro-2-methoxy-, (17.beta.)- (9CI)

(CA INDEX NAME)

OTHER NAMES:

CN 4-Fluoro-2-methoxyestradiol

FS STEREOSEARCH

MF C19 H25 F O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 105:146466

REFERENCE 2: 103:65176

L72 ANSWER 29 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 94761-90-7 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-methoxy-4-methyl- (7CI) (CA INDEX NAME)

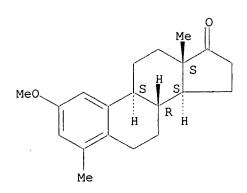
FS STEREOSEARCH

MF C20 H26 O2

LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L72 ANSWER 30 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 94440-60-5 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(hydroxymethyl)-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1, 3, 5(10) -triene-3, 17. beta. -diol, 2-(hydroxymethyl) - (7CI)

FS STEREOSEARCH

MF C19 H26 O3

LC STN Files: CA, CAOLD, CAPLUS, TOXCENTER, TOXLIT

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 136:31870

REFERENCE 2: 130:262306

REFERENCE 3: 125:105405

REFERENCE 4: 124:289981

REFERENCE 5: 122:187857

REFERENCE 6: 120:192069

L72 ANSWER 31 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 77736-30-2 REGISTRY

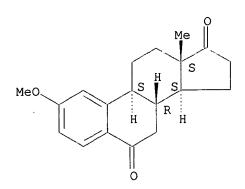
CN Estra-1, 3, 5(10) - triene-6, 17-dione, 2-methoxy- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H22 O3

LC STN Files: CA, CAPLUS

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 95:7573

L72 ANSWER 32 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 77736-29-9 REGISTRY

CN Estra-1,3,5(10)-trien-6-one, 17-hydroxy-2-methoxy-, (17.beta.)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H24 O3 LC STN Files:

CA, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 95:7573

L72 ANSWER 33 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 77736-24-4 REGISTRY

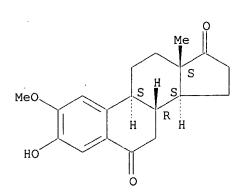
CN Estra-1,3,5(10)-triene-6,17-dione, 3-hydroxy-2-methoxy- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H22 O4

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 97:163313

REFERENCE 2: 95:7573

L72 ANSWER 34 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 69833-94-9 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-methoxy-, O-methyloxime (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H27 N O3

LC STN Files: SPECINFO

Absolute stereochemistry.

Double bond geometry unknown.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L72 ANSWER 35 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 69540-63-2 REGISTRY

CN Estra-1, 3, 5(10) -trien-6-one, 3, 17-dihydroxy-2-methoxy-, (17.beta.)- (9CI)

(CA INDEX NAME)

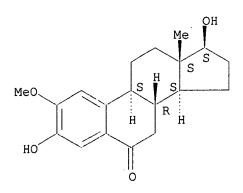
FS STEREOSEARCH

DR 77736-21-1

MF C19 H24 O4

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 134:295993

REFERENCE 2: 97:163313

REFERENCE 3: 95:7573

REFERENCE 4: 92:144013

REFERENCE 5: 92:108010

REFERENCE 6: 90:121859

L72 ANSWER 36 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 69505-03-9 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-methoxy-, (.+-.)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H24 O2

LC STN Files: BEILSTEIN*, CA, CAPLUS, GMELIN*

(*File contains numerically searchable property data)

Relative stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 93:114810

REFERENCE 2: 90:121861

L72 ANSWER 37 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 65968-97-0 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 4, 17-triol, 2-methoxy-, (17.beta.) - (9CI) (CA

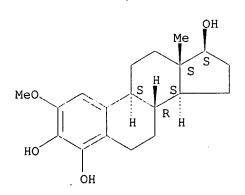
INDEX NAME)

FS STEREOSEARCH

MF C19 H26 O4

LC STN Files: CA, CAPLUS

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 88:136839

L72 ANSWER 38 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 65968-96-9 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3,4-dihydroxy-2-(phenylmethoxy)- (9CI) (CA

INDEX NAME)

STN Files:

FS STEREOSEARCH

MF C25 H28 O4

LC

CA, CAPLUS

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 88:136839

L72 ANSWER 39 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 59939-20-7 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-methoxy-, hydrogen sulfate,

(17.beta.) - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 O3 . x H2 O4 S

LC STN Files: CA, CAPLUS

CM 1

CRN 7664-93-9

CMF H2 O4 S

CM 2

CRN 362-07-2

CMF C19 H26 O3

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 84:148461

L72 ANSWER 40 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 58562-34-8 REGISTRY

CN Estra-1, 3, 5(10) -trien-17-one, 3, 4-dihydroxy-2-methoxy- (9CI) (CA INDEX

NAME)

OTHER NAMES:

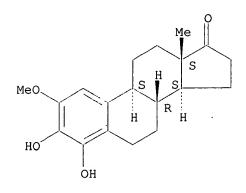
CN 2,3,4-Trihydroxy-1,3,5(10)-estratrien-17-one 2-methyl ether

FS STEREOSEARCH

MF C19 H24 O4

LC STN Files: CA, CAPLUS

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 88:136839

REFERENCE 2: 84:119044

REFERENCE 3: 84:102907

L72 ANSWER 41 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 52691-63-1 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(1-methyl-2-propenyl)-,

[2(S), 17.beta.] - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C22 H30 O2

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 80:146402

L72 ANSWER 42 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 52619-49-5 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-(1-methyl-2-propenyl)-, (17.beta.)-(9CI) (CA INDEX NAME)

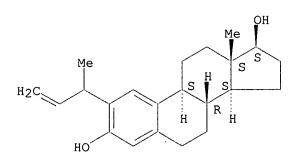
FS STEREOSEARCH

MF C22 H30 O2

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 80:146402

L72 ANSWER 43 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 51933-36-9 REGISTRY

CN Estra-1, 3, 5(10) -trien-17-one, 2-(2-chloro-2-propenyl)-3-hydroxy- (9CI)

(CA INDEX NAME)

FS STEREOSEARCH MF C21 H25 C1 O2

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 81:91784

L72 ANSWER 44 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 42028-20-6 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 3-hydroxy-2-methoxy-4-methyl- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Methoxy-4-methylestrone

FS STEREOSEARCH

MF C20 H26 O3

LC STN Files: CA, CAPLUS, TOXLIT

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 79:27594

L72 ANSWER 45 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 42028-17-1 REGISTRY

CN Estra-1,3,5(10)-triene-1,17-diol, 2-methoxy-4-methyl-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-triene-1,17.beta.-diol, 2-methoxy-4-methyl- (7CI)

FS STEREOSEARCH

MF C20 H28 O3

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, TOXLIT

(*File contains numerically searchable property data)

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 79:27594

L72 ANSWER 46 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 41259-43-2 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-azido-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Azidoestradiol

FS STEREOSEARCH

MF C18 H23 N3 O2

LC STN Files: BIOSIS, CA, CAPLUS, TOXCENTER, TOXLIT

Absolute stereochemistry.

7 REFERENCES IN FILE CA (1967 TO DATE)
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 118:96907

REFERENCE 2: 101:123208

REFERENCE 3: 96:116050

REFERENCE 4: 94:99013

REFERENCE 5: 91:168869

REFERENCE 6: 80:78582

REFERENCE 7: 79:137346

L72 ANSWER 47 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 41164-50-5 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-azido-3-hydroxy- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-Azidoestrone

FS STEREOSEARCH

MF C18 H21 N3 O2

LC STN Files: CA, CAPLUS, TOXCENTER, TOXLIT

Absolute stereochemistry.

6 REFERENCES IN FILE CA (1967 TO DATE)

6 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 118:96907

REFERENCE 2: 94:99013

REFERENCE 3: 92:53642

REFERENCE 4: 91:168869

REFERENCE 5: 80:78582

REFERENCE 6: 79:137346

L72 ANSWER 48 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 35577-55-0 REGISTRY

CN Estra-1,3,5(10)-trien-17-ol, 2-methoxy-4-methyl-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-trien-17.beta.-ol, 2-methoxy-4-methyl- (6CI, 7CI)

OTHER NAMES:

CN 2-Methoxy-4-methylestratriene-17.beta.-ol

FS STEREOSEARCH

MF C20 H28 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, TOXLIT

(*File contains numerically searchable property data)

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 79:27594

REFERENCE 2: 76:113417

L72 ANSWER 49 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 31559-55-4 REGISTRY

CN 19-Nor-17.alpha.-pregna-1,3,5(10)-triene-2-carbonitrile, 3,17-dihydroxy-

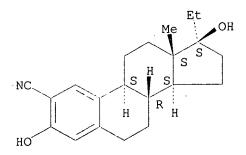
(8CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H27 N O2

LC STN Files: CA, CAPLUS, IFICDB, IFIPAT, IFIUDB

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 74:112314

L72 ANSWER 50 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 27293-73-8 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-methoxy- (8CI, 9CI) (CA INDEX NAME)

FS STEREOSEARCH

DR 27293-89-6

MF C19 H24 O2

LC STN Files: BEILSTEIN*, GMELIN*, TOXLIT

(*File contains numerically searchable property data)

L72 ANSWER 51 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 22145-26-2 REGISTRY

CN Estra-1, 3, 5(10) -triene-3, 17-diol, 2-(phenylmethoxy)-, (17.beta.)- (9CI)

(CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-triene-3,17.beta.-diol, 2-(benzyloxy)- (8CI)

FS STEREOSEARCH

MF C25 H30 O3

LC STN Files: CA, CAPLUS, ÜSPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 124:317590

REFERENCE 2: 119:117610

REFERENCE 3: 77:101989

REFERENCE 4: 70:106759

L72 ANSWER 52 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 17553-16-1 REGISTRY

CN Estra-1,3,5(10)-trien-17.beta.-ol, 2-methoxy- (6CI, 7CI, 8CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C19 H26 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS

(*File contains numerically searchable property data)

3 REFERENCES IN FILE CA (1967 TO DATE)

3 REFERENCES IN FILE CAPLUS (1967 TO DATE)

3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 73:116788

REFERENCE 2: 72:121765

REFERENCE 3: 68:69179

L72 ANSWER 53 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 16373-38-9 REGISTRY

CN Estra-1,3,5(10)-triene-3,17.beta.-diol, 2-methoxy-1-methyl- (8CI) (CA

INDEX NAME)

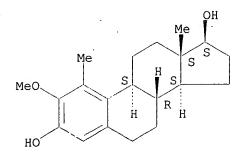
FS STEREOSEARCH

MF C20 H28 O3

LC STN Files: BEILSTEIN*, CA, CAPLUS

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 67:117038

L72 ANSWER 54 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 10506-80-6 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-allyl-3-hydroxy- (7CI, 8CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estrone, 2-allyl- (6CI)

FS STEREOSEARCH

MF C21 H26 O2

LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

5 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L72 ANSWER 55 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 10506-67-9 REGISTRY

CN Estra-1,3,5(10)-triene-3,17.beta.-diol, 2-allyl- (7CI, 8CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C21 H28 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, TOXLIT (*File contains numerically searchable property data)

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 66:102215

L72 ANSWER 56 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 10332-20-4 REGISTRY

CN Estra-1,3,5(10)-triene-3,17-diol, 2-methoxy-, (17.alpha.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-triene-3,17.alpha.-diol, 2-methoxy- (8CI)

FS STEREOSEARCH

MF C19 H26 O3

LC STN Files: BEILSTEIN*, CA, CAPLUS, TOXLIT

(*File contains numerically searchable property data)

2 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

REFERENCE 1: 110:166220

REFERENCE 2: 105:146466

L72 ANSWER 57 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 7683-30-9 REGISTRY

CN 17H-Cyclopenta[a]phenanthren-17-one, 6,7,8,9,11,12,13,14,15,16-decahydro-3-

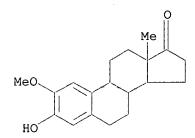
hydroxy-2-methoxy-13-methyl- (8CI) (CA INDEX NAME)

FS 3D CONCORD

MF C19 H24 O3

LC STN Files: BEILSTEIN*

(*File contains numerically searchable property data)



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L72 ANSWER 58 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 6564-02-9 REGISTRY

CN Estra-1,3,5(10)-triene-2-carbonitrile, 3-hydroxy-17-oxo- (7CI, 8CI, 9CI)

(CA INDEX NAME)

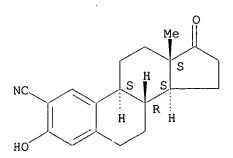
FS STEREOSEARCH

MF C19 H21 N O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, IFICDB, IFIPAT, IFIUDB

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 113:191714

REFERENCE 2: 74:112314

L72 ANSWER 59 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 6562-00-1 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2-(1,1-dimethylallyl)-3-hydroxy- (7CI, 8CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C23 H30 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS
(*File contains numerically searchable property data)

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 70:68602

L72 ANSWER 60 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 5976-42-1 REGISTRY

CN Estra-1,3,5(10)-triene-2-carbonitrile, 3,17-dihydroxy-, (17.beta.)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estra-1,3,5(10)-triene-2-carbonitrile, 3,17.beta.-dihydroxy- (7CI, 8CI)

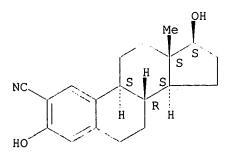
FS STEREOSEARCH

MF C19 H23 N O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, CASREACT, IFICDB, IFIPAT, IFIUDB

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 113:191714

REFERENCE 2: 74:112314

L72 ANSWER 61 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 4967-94-6 REGISTRY

CN Estra-1,3,5(10)-triene, 2-methoxy- (8CI) (CA INDEX NAME)

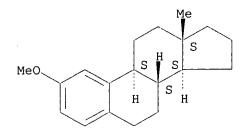
FS STEREOSEARCH

MF C19 H26 O

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1967 TO DATE)

2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 73:116788

REFERENCE 2: 70:78228

L72 ANSWER 62 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 4953-96-2 REGISTRY

CN Estra-1,3,5(10)-trien-3-ol, 2-methoxy- (7CI, 8CI) (CA INDEX NAME)

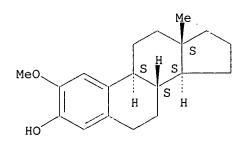
FS STEREOSEARCH

MF C19 H26 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS

(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 70:78228

L72 ANSWER 63 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 2120-13-0 REGISTRY

CN Estra-1,3,5(10)-triene-3,17.beta.-diol, 2,4-diallyl- (7CI, 8CI) (CA INDEX

FS STEREOSEARCH

MF C24 H32 O2

LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, TOXLIT (*File contains numerically searchable property data)

Absolute stereochemistry.

$$H_2C$$
 H_2C
 H_2C
 H_2C

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 66:102215

L72 ANSWER 64 OF 64 REGISTRY COPYRIGHT 2002 ACS

RN 2120-12-9 REGISTRY

CN Estra-1,3,5(10)-trien-17-one, 2,4-diallyl-3-hydroxy- (7CI, 8CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Estrone, 2,4-diallyl- (6CI)

FS STEREOSEARCH

MF C24 H30 O2

LC STN Files: BEILSTEIN*, CAOLD

(*File contains numerically searchable property data)

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

5 REFERENCES IN FILE CAOLD (PRIOR TO 1967)